

HepaCAM-PIK3CA axis regulates the reprogramming of glutamine metabolism to inhibit prostate cancer cell proliferation

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Abstract. Energy metabolism reprogramming is becoming an increasingly important hallmark of cancer. Specifically, cancers tend to undergo metabolic reprogramming to upregulate a cell-dependent glutamine (Gln) metabolism. Notably, hepatocellular cell adhesion molecule (HepaCAM) has been previously reported to serve a key role as a tumour suppressor. However, the possible regulatory role of HepaCAM in Gln metabolism in prostate cancer (PCa) remains poorly understood. In the present study, bioinformatics analysis predicted a significant negative correlation among the expression of HepaCAM, phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit α (PIK3CA), glutaminase (GLS) and solute carrier family 1 member 5 (SLC1A5), components of Gln metabolism, in clinical and genomic datasets. Immunohistochemistry results verified a negative correlation between HepaCAM and PIK3CA expression in PCa tissues. Subsequently, liquid chromatography-tandem mass spectrometry (LC-MS/MS) and gas chromatography-mass spectrometry (GC-MS) assays were performed, and the results revealed markedly reduced levels of Gln and metabolic flux in the blood samples of patients with PCa and in PCa cells. Mechanistically, overexpression of HepaCAM inhibited Gln metabolism and proliferation by regulating PIK3CA in PCa cells. In addition, Gln metabolism was discovered to be stress-resistant in PCa cells, since the expression levels of GLS

and SLC1A5 remained high for a period of time after Gln starvation. However, overexpression of HepaCAM reversed this resistance to some extent. Additionally, alpelisib, a specific inhibitor of PIK3CA, effectively potentiated the inhibitory effects of HepaCAM overexpression on Gln metabolism and cell proliferation through mass spectrometry and CCK-8 experiments. In addition, the inhibitory effect of PIK3CA on the growth of tumor tissue in nude mice was also confirmed by immunohistochemistry *in vivo*. To conclude, the results from the present study revealed an abnormal Gln metabolic profile in the blood samples of patients with PCa, suggesting that it can be applied as a clinical diagnostic tool for PCa. Additionally, a key role of the HepaCAM/PIK3CA axis in regulating Gln metabolism, cell proliferation and tumour growth was identified. The combination of alpelisib treatment with the upregulation of HepaCAM expression may serve as a novel method for treating patients with PCa.

Introduction

Prostate cancer (PCa) is one of the most prevalent urological malignancies, with the second-highest rate of cancer-associated mortality among the male population in Western countries (1). Over the past decade, with the rapid growth of an aging population in China, the incidence, morbidity pattern and mortality rates of PCa have all been increasing on an annual basis. Notably, the majority of patients with PCa develop incurable disease due to advanced regional disease and/or distant metastasis (2).

Cancer cells exhibit the ability to survive and proliferate even under unfavourable conditions, such as nutrient-poor environments (3). Previous reports have demonstrated that energy metabolism reprogramming, which potentiates cell proliferation by adjusting energy metabolism, is an emerging hallmark of cancer (4,5). In particular, metabolic reprogramming towards glutamine (Gln) and cell-dependent Gln metabolic activity have both been found to be increased in a variety of cancers and this is a common metabolic alteration in cancers, where the importance of Gln as a nutrient is considered second only to glucose (5). Gln is one of the most abundant free amino acids and can regulate ATP production, macromolecular synthesis and signal transmission in cancer

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cells by donating its nitrogen and carbon atoms (6). In general, Gln is imported into the cell through specific transporters, such as solute carrier family 1 member 5 (SLC1A5; also known as ASCT2). Gln metabolism begins with its conversion to glutamate (Glu), in a process catalysed by glutaminase (GLS), with type I glutaminase (GLS1) as the dominant isoform expressed in humans (7,8). Additional studies have revealed a close relationship between Gln metabolism and PCa, such that Gln has been proposed to be a potential marker for the diagnosis or treatment of PCa (9-12).

Hepatocellular cell adhesion molecule (HepaCAM) is a novel type of immunoglobulin (Ig) cell adhesion molecule that has shown tumour suppressor-like characteristics and was first isolated from the human liver (13). Under physiological conditions, HepaCAM is expressed at high levels in normal tissues and cells (14). However, the expression of HepaCAM was previously found to be downregulated or even lost in several tumour tissues and cells, including PCa (13-15). In addition, HepaCAM expression has been negatively associated with the proliferation, invasion and the Warburg effect in malignancies of the urinary system, including bladder cancer and renal cell carcinoma (16-19). Overexpression of HepaCAM has also been found to inhibit the proliferation and invasion of PCa cells (20). However, the possible regulatory mechanism of HepaCAM in regards to Gln metabolism in PCa remains poorly understood.

Previous reports have found the PI3K signalling pathway to be one of the most frequently dysregulated pathways in cancer (21-24). Phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit α (PIK3CA) is one of the most common oncogenes which encodes the p110 α catalytic subunit of PI3K (25). PIK3CA can be constitutively activated by two main different mechanisms, namely activating mutation and amplification (26). Accumulating evidence has revealed that PIK3CA is associated with Gln metabolism reprogramming in numerous types of cancers (27-30). Consequently, PIK3CA may also be a key regulatory target for reprogramming Gln metabolism in PCa.

The present study explored the possible role of HepaCAM in Gln metabolism in PCa, in addition to attempting the identification of the potential relationship between HepaCAM and PIK3CA expression. It was hypothesized that the overexpression of HepaCAM can inhibit Gln metabolism reprogramming and PCa cell proliferation by regulating PIK3CA.

Materials and methods

Bioinformatics analysis. A total of 19 PCa datasets (source link: https://www.cbioportal.org/study/summary?id=prad_mich%2Cprad_su2c_2019%2Cprad_su2c_2015%2Cprad_mespc_mskcc_2020%2Cprad_broad_2013%2Cprad_broad%2Cprad_cpcg_2017%2Cprad_cdk12_mskcc_2020%2Cprad_mskcc%2Cprad_p1000%2Cprad_eurrol_2017%2Cprad_tcga_pan_can_atlas_2018%2Cprad_mskcc_chenyl_organoids_2014%2Cprad_mskcc_2017%2Cprad_msk_2019%2Cprad_tcga_pub%2Cprad_tcga%2Cprostate_dkfz_2018%2Cmpcproject_broad_2021), including The Cancer Genome Atlas, TCGA (<https://www.cancer.gov/about-nci/organization/ccg/research/structural-genomics/tcga>) were downloaded from the cBioPortal (<https://www.cbioportal.org>). In total, 7 datasets had HepaCAM

structural variation, mutation or copy number variation (CNA) data, with 2,093 cases (94.19%) belonging to PCa. In addition, 19 datasets had PIK3CA structural variation, mutation or CNA data, including 6,671 (95.5%) cases of PCa (including castration-resistant PCa and prostate neuroendocrine carcinoma) and 324 cases (4.5%) of a normal group (N). The present study was conducted in accordance with the Declaration of Helsinki (as revised in 2013).

Blood and tissue samples. Between 2016 and 2018, tissue specimens were collected from 17 patients (age range, 53-86 years; mean age, 70) with benign prostatic hyperplasia (BPH) and 44 patients (age range, 55-91 years; mean age, 71) with PCa. In addition, blood specimens were collected from 43 individuals containing normal samples and 66 patients with PCa from 2018 to 2019. All patients were recruited from the Department of Urology, The First Affiliated Hospital of Chongqing Medical University (Chongqing China). All samples were verified by a pathologist, who histologically diagnosed the samples as either BPH or PCa. The present study was approved by the Ethics Committee of Chongqing University. All patients provided written informed consent prior to specimen acquisition.

Immunohistochemistry. All samples involved in the experiment, including human PCa and BPH specimens, were cut into paraffin sections. Detection of HepaCAM and PIK3CA expression in PCa and BPH tissue samples was performed using the IHC staining procedure as described previously (31). The primary antibodies used were as follows: anti-HepaCAM (cat. no. MAB4108; 1:150; ProteinTech Group, Inc.) and anti-PIK3CA (cat. no. 4249T; 1:200; Cell Signaling Technology, Inc.). Tissues underwent fixation with 4% paraformaldehyde, were embedded in paraffin and then subjected to standard dewaxing and rehydration. The sections were incubated in citric acid buffer (pH 6.0) for 15 min for antigen retrieval, followed by incubation for 10 min with 3% H₂O₂ solution to inactivate endogenous enzymatic activity. The sections were then incubated with anti-HepaCAM antibody and anti-PIK3CA antibody for 1 h at 25°C followed by Elivision Plus Polymer HRP (mouse/rabbit) immunohistochemistry kit (cat. no. KIT-9903; 1:1,000; Maxim Biotech, Inc.) for 30 min at 20°C. The results of HepaCAM and PIK3CA expression in PCa and BPH tissue samples was analysed according to a previously described procedure (31).

Cell culture and treatment. Prostate cell lines PC3 (cat. no. ZQ0041), LNCaP (cat. no. ZQ0039) and RWPE-1 (cat. no. ZQ0351) were obtained from Shanghai ZhongQiao XinZhou Biotechnology Co., Ltd. RWPE-1 cells were cultured with Keratinocyte-SFM medium (cat. no. 10744-019; Gibco; Thermo Fisher Scientific, Inc.), whereas cancer cell lines PC3 and LNCaP cells were cultured with RPMI-1640 media (Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% FBS (Gibco; Thermo Fisher Scientific, Inc.) and 100 U/ml penicillin/streptomycin (Beyotime Institute of Biotechnology). All cells were cultured in a 37°C incubator containing 5% CO₂ and 1% O₂ with 45-65% humidity. Adenoviruses encoding HepaCAM (Ad-HepaCAM) were stored at -80°C and amplified in 293A cells. The viral fluid was obtained after the repeated freezing and thawing of the 293A cells. The prostate cancer cells were transfected with either Ad-HepaCAM or Ad-GFP,

respectively. Cells (1×10^5 /well) were seeded into 6-well plates and incubated with 2 ml complete medium for 12 h, before being transfected with adenoviruses and/or alpelisib ($8 \mu\text{M}$, 24 h; MedChemExpress, USA). After 24 or 72 h of incubation, subsequent experiments were performed.

Gln deprivation experiments. After the cells had completely attached to the culture dish, the RPMI-1640 medium was removed and PC3 and LNCaP cells were washed once with PBS. Gln-free medium was prepared using RPMI-1640 devoid of Gln (cat. no. A14431; Thermo Fisher Scientific, Inc.) but other nutrients were retained at the same concentrations as those in the standard complete RPMI-1640 media. Complete RPMI-1640 medium was prepared by supplementing with 2 mM L-Gln (Beijing Solarbio Science & Technology Co., Ltd.). All media were supplemented with 10% dialyzed FBS (Gemini Bio Products) and 100 U/ml penicillin/streptomycin. All cells were cultured in a 37°C incubator containing 5% CO_2 and 1% O_2 with 45–65% humidity. The incubation time depended on the experimental requirements (24–72 h).

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). RNA was extracted from RWPE-1, PC3 and LNCaP cells using TRIzol (Thermo Fisher Scientific, Inc.), and PrimeScript RT reagent kits (Takara Bio, Inc.) were used to reverse transcribe the RNA. Real-time PCR was performed with the SYBR Premix Ex Taq™ II kit (cat. no. RR820A; TaKaRa, Inc.). Notably, β -actin was used as the standard reference. The thermocycling conditions for qPCR were as follows: initial denaturation at 95°C for 3 min; followed by 40 cycles of 95°C for 10 sec, 60°C for 20 sec and 72°C for 20 sec; and final extension at 72°C for 5 min. The comparative $2^{-\Delta\Delta\text{C}_q}$ method was used to calculate the relative expression level of mRNA (32). Each experiment was repeated ≥ 3 times. The primer sequences (5'-3') used were as follows: *HepaCAM* forward, TGTACAGCTGCATGGTGGAGA and reverse, TCTGGTTTCAGGCGGTCATCA; *PIK3CA* forward, CCACGACCATCATCAGGTGAA and reverse, CCTCACGGAGGCATTCTAAAGT; *GLS* forward, TACTGAGCCCTGAAGC and reverse, TCCAGAGGAGGAGACC; *SLC1A5* forward, GAGCTGCTTATCCGCTTCTTC and reverse, GGGGCGTACCACATGATCC; *Cyclin D1* forward, GCTGGAGCCCGTGAAAAAGA and reverse, CTCCGCCTCTGGCATTTTG; proliferating cell nuclear antigen (*PCNA*) forward, TCAAGAAGGTGTTGGAGGCA and reverse, CAGCGGTAGGTGTCGAAGC and β -actin forward, TGACGTGGACATCCGCAAAG and reverse, CTGGAAGGTGGACAGCGAGG.

Western blotting. Total proteins were extracted by RIPA buffer (cat. no. 9806; Cell Signaling Technology, Inc.) containing the protease inhibitor phenyl methane sulfonyl fluoride and phosphatase inhibitors (33,34). Nuclear and plasma proteins were extracted using the nuclear and cytoplasmic protein extraction reagent (Beyotime Institute of Biotechnology). The BCA kit (Beyotime Institute of Biotechnology) was used to measure protein concentration. SDS-PAGE (10% gel) was used for protein separation ($50 \mu\text{g}$). The following antibodies were used: anti-HepaCAM (cat. no. 18177-1-AP; 1:1,000; ProteinTech Group, Inc.), anti-PIK3CA (cat. no. 4249T; 1:1,000; Cell Signaling Technology, Inc.), anti-AKT1

(cat. no. abs131788; 1:1,000; Absin Bioscience, Inc.), anti-phosphorylated (p-)AKT1 (Ser473; cat. no. abs130002; 1:1,000; Absin Bioscience, Inc.), anti-p-AKT1 (Thr308; cat. no. abs130889; 1:1,000; Absin Bioscience, Inc.), anti-GLS1 (cat. no. 56750; 1:1,000, Cell Signaling Technology, Inc.), anti-SLC1A5 (cat. no. 20350-1-AP; 1:3,000; ProteinTech Group, Inc.), anti-cyclin D1 (cat. no. 55506; 1:500, Cell Signaling Technology, Inc.), anti-PCNA (cat. no. 13110; 1:500; Cell Signaling Technology, Inc.), anti- β -actin (cat. no. 20536-1-AP; 1:5,000; ProteinTech Group, Inc.), and HRP-conjugated anti-rabbit-IgGk binding protein secondary antibody (cat. no. ab205718; 1:5,000; Abcam). Enhanced Chemiluminescence Detection Kit was utilized for the visualization of protein bands (Suzhou New Saimei Biotechnology Co., Ltd.).

Cell Counting Kit-8 (CCK-8) assay. PC3 and LNCaP cells (3×10^3 /well) were first cultured in 96-well plates overnight before being transfected/treated with Ad-HepaCAM/-Gln/alpelisib or control. After 0, 24, 48 or 72 h, $10 \mu\text{l}$ CCK-8 reagent (cat. no. 354550 Shanghai Univ Biotechnology Co., Ltd.) was added and incubated for 1–4 h at 37°C . Finally, absorbance in each well was measured at 450 nm. A total of four wells were used for each treatment group per experiment.

Colony formation assay. PC3 and LNCaP cells (500/well) following blank treatment and administration of Ad-GFP or Ad-HepaCAM were added into 6-well plates and cultured in a 37°C incubator containing 5% CO_2 and 1% O_2 with 45–65% humidity. for 7–14 days. The cells were then fixed with methanol for 2 min and stained with 0.5% crystal violet for 10 min at room temperature and photographed for cell counting. The images of each well were scanned using an Epson scanner GT-X970 (Seiko Epson Corp.), and the colonies were counted using ImageJ 1.52v software (National Institutes of Health). Each treatment group used three wells per experiment and each experiment was repeated three times independently.

Flow cytometry. PC3 and LNCaP cells were transferred into 6-well plates overnight and transfected/treated with Ad-HepaCAM or alpelisib. After 72 h, the cells were washed two to three times with PBS, before being digested using 0.25% trypsin, washed twice again with PBS and transferred into Eppendorf tubes. The cells were counted and 1×10^6 cells were resuspended with $100 \mu\text{l}$ PBS before $500 \mu\text{l}$ cooled 75% ethanol was added after centrifugation. Cell cycle distribution was detected using the CytoFLEX Flow Cytometer (Beckman Coulter, Inc.) and the results were analysed using ModFitLT 5 software (Verity Software House, Inc.).

Immunofluorescence. PC3 and LNCaP cells (3×10^4) were cultured on coverslips in 24-well plates following administration of Ad-HepaCAM or/and alpelisib, and then the cells were incubated for 72 h. The cells were then fixed with 4% paraformaldehyde, 0.1–1% Triton X-100 and 5% goat blocking serum in sequence as previously described (35). Primary antibodies against anti-Ki67 (cat. no. ab197234; 1:100, Abcam), secondary antibodies and 4,6-diamidino-2-phenyl-indole (Zhongshan Golden Bridge Biotechnology, Ltd.; OriGene Technologies, Inc.) were used for incubation with the cells.

Fluorescence images were captured using a confocal laser scanning microscope (Nikon Corp.) at x400 magnification. The Ki-67-positive index was analysed by Image J (v4.6.2, National Institutes of Health).

Liquid chromatography-tandem mass spectrometry (LC-MS/MS) and gas chromatography-mass spectrometry (GC-MS) assay. The trypsin-digested PC3 and LNCaP cells ($>1 \times 10^7$) from the different treatment groups were collected, before the supernatant was discarded after centrifugation. Cells were washed three times with PBS before being fully disrupted using a sonicator (16,000 x g). Finally, pre-cooled methanol was added for fixation. After fixation, the lysed cells were suspended on a dried blood spot (DBS) to be sufficiently saturated. Fresh blood samples from different patients were collected, prepared in a DBS format on a filter paper and stored at 4°C. MS/MS analysis was achieved using a QTRAP triple quadrupole mass spectrometer (AB SCIEX 3200; AB SCIEX, USA) coupled to a LC system (Shimadzu 20ADXR; Shimadzu).

Approximately 20 μ l of working solution was directly injected for analysis without prior chromatographic separation. A mixed liquor (acetonitrile:H₂O=1:1) containing 0.05% FA was used as the mobile phase. The mass spectrometer was equipped with an electrospray ionization (ESI) and operated in the positive ion mode to monitor the m/z transitions for all peptides and IS. For each peptide, two multiple reaction monitorings (MRMs) were simultaneously monitored. A scheduled MRM acquisition method was constructed using manually optimized declustering potentials (DP), collision energies (CE), collision cell entrance (CEP) and exit potentials (CXP). All chromatograms were analysed with Analyst 1.5 software (AB SCIEX, USA) using internal standards, and the peak-area ratios were applied for further calculations (36).

Xenograft model in vivo. A total of 10 male nude mice (4 weeks of age; body weight, 10–15 g) were purchased from the Shanghai SLAC Laboratory Animal Co., Ltd. All rats were housed at 23–25°C with 50–60% humidity, 12-h light/dark cycle and food and water *ad libitum*. The present study was performed in accordance with the Animal Research Advisory Committee (ARAC) Guidelines (<https://oacu.oir.nih.gov/animal-research-advisory-committee-guidelines>) and was approved by the Ethics Committee of Chongqing Medical University (Chongqing, China). A prostate cancer xenograft model was established by the subcutaneous injection of LNCaP cells into the right flank of 10 mice (5×10^6 cell per mouse). These nude mice were then randomly divided into a dosing group and control group. Alpelisib was dissolved in 0.5% sodium carboxyl methyl cellulose (CMC-NA). Alpelisib (12.5 mg/kg) was administered daily through an oral gavage for 5 consecutive days, and then 1 day off and the above process was repeated once. The same amount of 0.5% CMC-NA was given by intragastric administration to the control group. Tumour size was assessed every day. After 4 weeks, the 10 mice were sacrificed by CO₂ euthanasia. The criteria for determining the end point of this animal experiment were as follows: i) the animal was dying or unable to move, or had no response after giving gentle stimulation; ii) difficulty breathing, with typical symptoms being salivation and/or cyanosis in the mouth and nose; iii) diarrhea or urinary incontinence; iv) the body weight was reduced by 20% of the body

weight during the experiment; v) inability to eat or drink; vi) the animal was showing clear signs of anxiety, irritability; vii) the weight of the tumour reached >10% of the animal's own body weight; viii) animal skin damage area accounted for >30% of the whole body; and ix) or infection and suppuration occurred. CO₂ euthanasia was performed using a CO₂ delivery system according to the ARAC guide (<https://oir.nih.gov/sourcebook/committees-advisory-ddir/animal-research-advisory-committee-arac>) for the experimental procedure. The mouse was first placed in the box full of air before the box was filled with a volume of 2 liters CO₂ to a volume of 10 liters at a constant rate (the volume displacement rate of CO₂ was 60% vol/min). Each mouse was checked for the lack of breathing and constriction of pupil size. After the breathing halted, the CO₂ flow was maintained for 1 min and the mouse was removed from the euthanasia box. Death was verified by the cessation of heartbeat and breathing, in addition to the reflexes disappearing. Tumour tissues were then isolated for comparing the size and weight before IHC experiments were performed. Tumour volume was calculated using the formula $V = \pi/6 \times L \times L \times W$, where V was the volume and L was the length and W was the width. The maximum volume and the maximum diameter of a single tumour presented in the present study were 764.5 mm³ and 12.1 mm, respectively.

Statistical analysis. SPSS (version 24.0; IBM Corp.) and GraphPad Prism (version 5; GraphPad Software, Inc.) were used for statistical analysis. Data are presented as the mean \pm SD. All experiments were repeated at least three times. Comparisons between two groups, other than comparisons between tumour and adjacent non-tumour samples, were analysed using the unpaired Student's t-test. A paired Student's t-test was used to compare HepaCAM and PIK3CA expression levels between tumour and adjacent non-tumour samples. Survival curves were assessed using log-rank tests. Comparisons among multiple groups were analysed using one-way ANOVA followed by Tukey's post hoc test. The χ^2 test was used to analyse the relationship between HepaCAM and PIK3CA expression or the concentration of Gln/Glu and each of the clinical parameters. The correlation between HepaCAM and PIK3CA expression in the tissue samples was evaluated using Spearman's rank correlation coefficient. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

Bioinformatics prediction of the correlation among HepaCAM, PIK3CA and Gln metabolism in clinical and genomic datasets. In a preliminary experiment, a correlation between the expression of HepaCAM and GLS was found in PCa cells, but the possibility of direct action was ruled out. The selection of PIK3CA for the present study was based on preliminary bioinformatics analysis, where protein interaction analysis in the STRING protein database found a possibility of protein interaction between HepaCAM and PIK3CA. In addition, previous studies have found that the overexpression of HepaCAM was able to inhibit the expression of c-Myc (37,38), which is an important regulatory target of SLC1A5 (39). Under these premises, the correlation among these components aforementioned was assessed through bioinformatics analysis. Comparative analysis of the different datasets in the

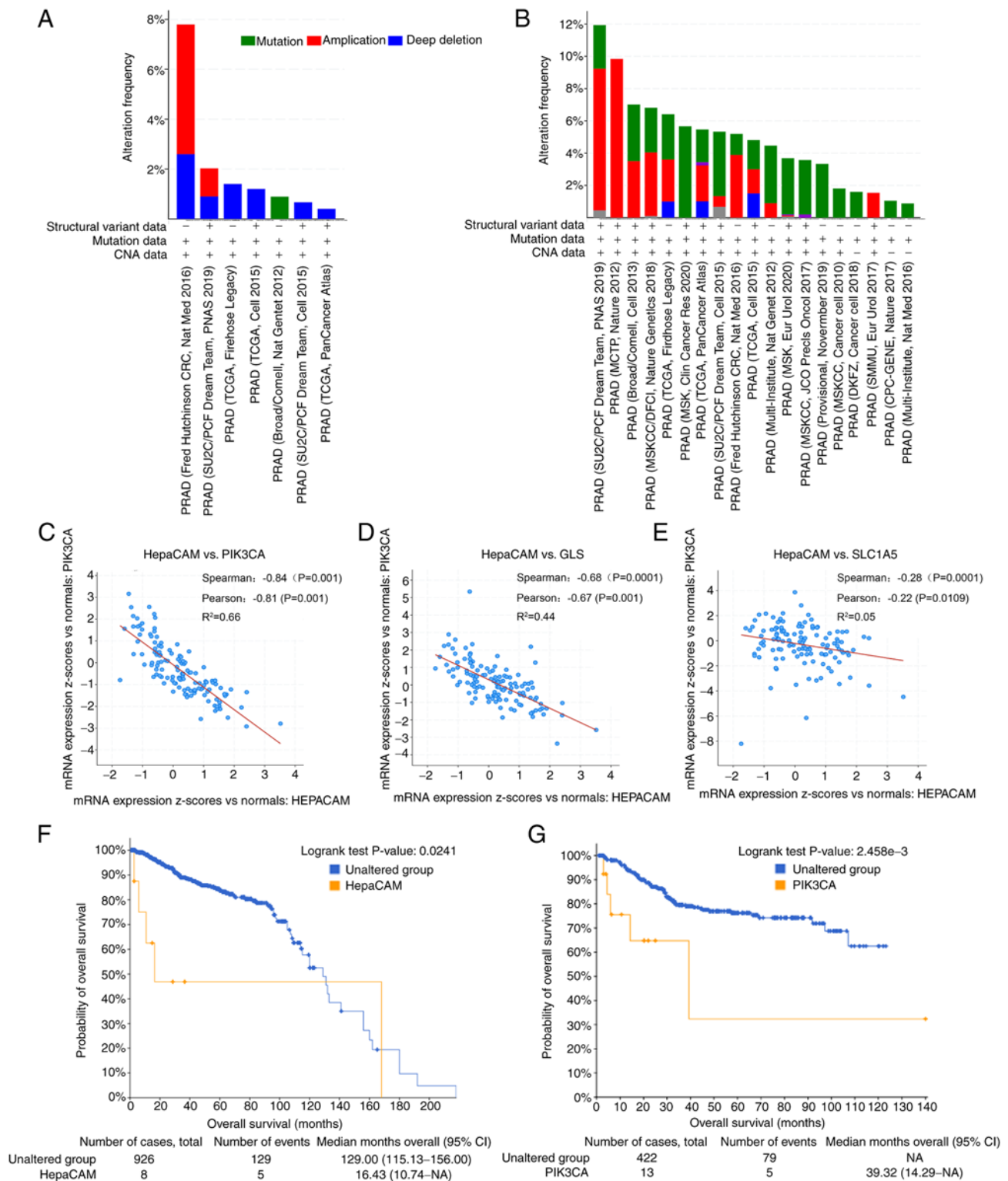


Figure 1. The correlation and difference between HepaCAM, PIK3CA, and glutamine metabolism-related molecules from the Bio-databases. (A and B) Variation types and proportion of HepaCAM and PIK3CA genes in different PCa datasets from the cBioportal. (C-E) RNA-seq mRNA expression data from GEO database (GSE21032) was used to compare the correlation of HepaCAM with PIK3CA, GLS and SLC1A5 in PCa tumors. (F and G) Overall survival curves of PCa patients according to HepaCAM and PIK3CA mRNA levels in TCGA. HepaCAM, hepatocellular cell adhesion molecule; PIK3CA, phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit α ; GLS, glutaminase; SLC1A5, solute carrier family 1 member 5. To note: the link to the 19 PCa datasets is provided in the *Bioinformatics analysis* section.

cBioportal database revealed little to no HepaCAM expression in the majority of cancer types tested (Fig. 1A). Furthermore, the main alteration in PIK3CA expression in PCa was found to be amplification (Fig. 1B), while mutation was the major

form of alteration in other solid tumours (40,41). Additionally, the co-expression analysis of genes in the databases found that the mRNA expression levels of *HepaCAM* exhibited a negative correlation with those of *PIK3CA* and components

Table I. Glutamine/glutamate concentrations and clinicopathological parameters of the PCa patients.

Characteristics	No. of specimens (%)	Glutamine (Gln) ($\mu\text{mol/l}$)			Glutamate (Glu) ($\mu\text{mol/l}$)		
		Median=9.766			Median=66.613		
		<9.766	≥ 9.766	P-value	<66.613	≥ 66.613	P-value
Histology							
Normal	43	0	43	0.001	41	2	0.001
PCa	66	47 (71)	19 (29)		13 (20)	53 (80)	
Age (years)							
<60	9 (14)	4 (6)	5 (8)	0.056	1 (2)	8 (12)	0.486
≥ 60	57 (86)	43 (65)	14 (21)		12 (18)	45 (68)	
PSA ($\mu\text{g/l}$)							
Median=20.67							
<20.67	31 (47)	25 (38)	6 (9)	0.233	4 (6)	27 (41)	0.192
≥ 20.67	35 (53)	22 (33)	13 (20)		9 (14)	26 (39)	
Gleason score of PCa							
≤ 6	19 (29)	10 (15)	9 (14)	0.014	5 (8)	14 (21)	0.690
7	23 (35)	15 (23)	8 (12)		4 (6)	19 (29)	
≥ 8	24 (36)	22 (33)	2 (3)		4 (6)	20 (30)	

Mann-Whitney test for 2 independent variables; Chi-square test for trend for the number of rows or columns >2. Numbers in bold font indicate statistical significance ($P < 0.05$). PCa, prostate cancer; PSA, prostate-specific antigen.

Table II. HepaCAM and PIK3CA in PCa tissues and clinicopathological parameters.

Characteristics	No. of specimens (%)	HepaCAM staining			PIK3CA staining		
		Positive	Negative	P-value	Positive	Negative	P-value
Total	44 (100)	15 (34)	29 (66)		38 (86)	6 (14)	
Age (year)							
<60	16 (36)	7 (16)	4 (20)	0.307	13 (29)	3 (7)	0.052
≥ 60	28 (64)	8 (18)	20 (46)		25 (57)	3 (7)	
Histological stage							
Ta-T2	29 (66)	10 (23)	19 (43)	0.939	27 (61)	2 (5)	0.07
T3-T4	15 (34)	5 (11)	10 (23)		11 (25)	4 (9)	
Gleason score							
<6	10 (23)	8 (18)	2 (5)	0.001	6 (14)	4 (9)	0.022
7	16 (36)	7 (16)	9 (20)		15 (34)	1 (2)	
≥ 8	18 (41)	0 (0)	18 (41)		17 (38)	1 (3)	

Mann-Whitney test for 2 independent variables; Chi-square test for trend for the number of rows or columns >2. Numbers in bold font indicate statistical significance ($P < 0.05$). HepaCAM, hepatocellular cell adhesion molecule; PCa, prostate cancer; PIK3CA, phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit α .

associated with Gln metabolism, *GLS* and *SLC1A5* ($P < 0.05$; Fig. 1C-E). Survival analyses of the datasets found that the patients whose samples possessed lower expression levels of HepaCAM displayed poorer overall survival ($P < 0.01$; Fig. 1F), whereas the survival trend for PIK3CA expression was opposite ($P < 0.05$; Fig. 1G).

Abnormal levels of Gln in blood samples and PCa cell lines. Blood samples were collected from 66 patients with PCa and 43 normal individuals for the detection of amino acid concentrations by LC-MS/MS and GC-MS testing. Among the 15 common amino acids detected, only the levels of Gln and arginine (Arg) in the blood samples of patients with PCa were

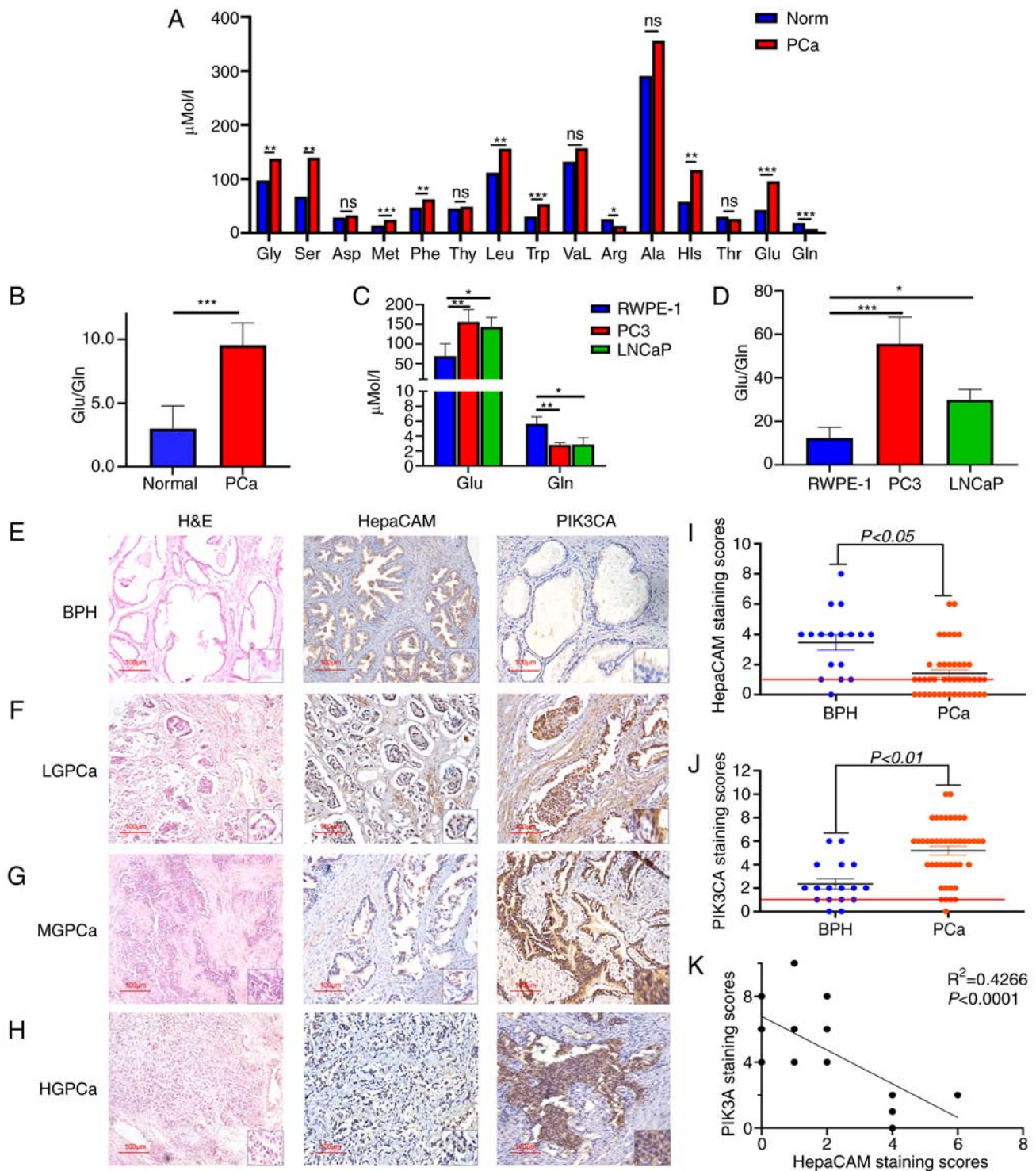


Figure 2. The significant difference in glutamine (Gln)/glutamate (Glu) levels in blood specimens or cells and HepaCAM and PIK3CA expression in tissues specimen. (A) Multiple common amino acid concentrations in different blood samples tested by mass spectrometry and analysed by Mann-Whitney test. * $P<0.05$, ** $P<0.01$, *** $P<0.001$; ns, not significant comparing PCa vs. normal (Norm) specimens. (B) Comparison of the ratio of Glu to Gln between PCa and normal samples. *** $P<0.001$ vs. normal specimens. (C and D) Difference in the concentration of Glu and Gln between prostate cancer cell lines PC3 and LNCaP and normal epithelial RWPE-1 cells. * $P<0.05$, ** $P<0.01$, *** $P<0.001$. (E-H) Representative hematoxylin and eosin (H&E) staining and IHC staining for HepaCAM and PIK3CA in PCa and BPH samples, Magnification, x200. (I and J) Staining scores of HepaCAM and PIK3CA expression in BPH and PCa tissues. (K) Correlation analysis between HepaCAM and PIK3CA in tissues as analysed by Spearman's correlation analysis. Glu, glutamate; Gln, glutamine; LGPCa, low-grade PCa; MGPCa, middle-grade PCa; HGPCa, high-grade PCa; BPH, benign prostatic hyperplasia; PCa, prostate cancer; HepaCAM, hepatocellular cell adhesion molecule; PIK3CA, phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit α .

significantly lower compared with those in the samples from the normal individuals ($P<0.05$; Fig. 2A; Table I). In addition, the concentration of Gln exhibited the most striking difference

among all other amino acids tested, where the main metabolic flow was towards the synthesis of glutamic acid ($P<0.001$; Fig. 2A and B; Table I).

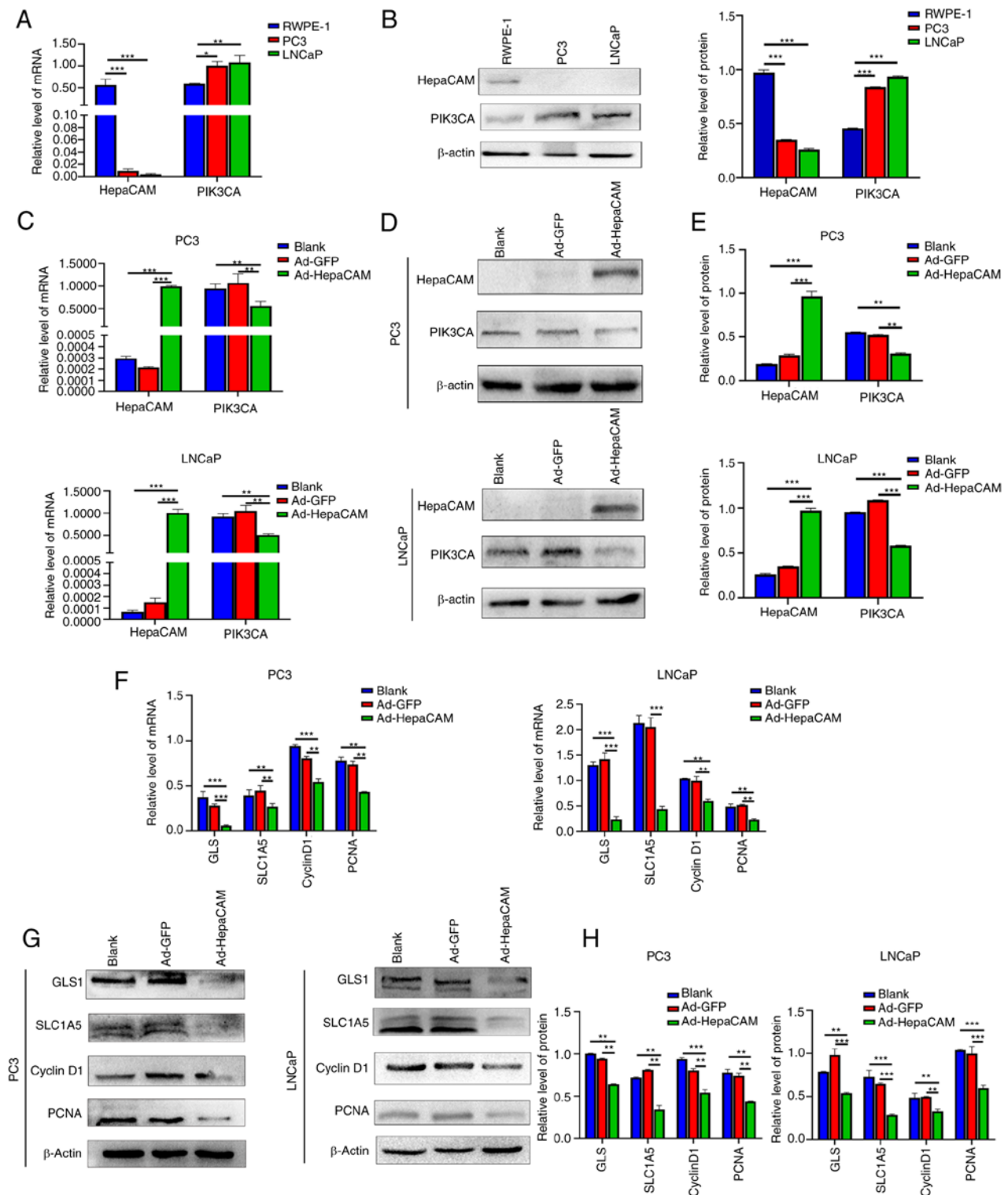


Figure 3. Overexpression of HepaCAM inhibits the expression of PIK3CA and glutamine metabolism or proliferation related molecules in PCa. (A and B) Expression of HepaCAM and PIK3CA in RWPE-1, PC3 and LNCaP cells was detected by (A) RT-qPCR and (B) western blot analysis. * $P < 0.05$, ** $P < 0.01$, and *** $P < 0.001$ compared with the RWPE-1 cells. (C-E) Overexpression of HepaCAM adenovirus and corresponding changes in PIK3CA at the (C) mRNA and (D and E) protein levels was verified in PC3 and LNCaP cells. (F-H) mRNA and protein levels of GLS, SLC1A5, cyclin D1 and PCNA in cells were detected by qPCR and western blot analysis after infection with adenoviral Ad-HepaCAM. β -actin was used as internal controls. Data are represented as mean \pm SD of 3 individual experiments. ** $P < 0.01$ and *** $P < 0.001$ vs. blank control. HepaCAM, hepatocellular cell adhesion molecule; PIK3CA, phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit α ; GLS, glutaminase; SLC1A5, solute carrier family 1 member 5; PCNA, proliferating cell nuclear antigen.

On a cellular level, the Gln levels in RWPE-1 cells was significantly higher compared with that in LNCaP and PC3 cells ($P < 0.01$; Fig. 2C), where the metabolic flux was also biased

towards synthesizing Glu ($P < 0.001$; Fig. 2D). Additionally, high levels of Gln in PCa were found to be significantly associated with the Gleason score (Table I). These results suggest

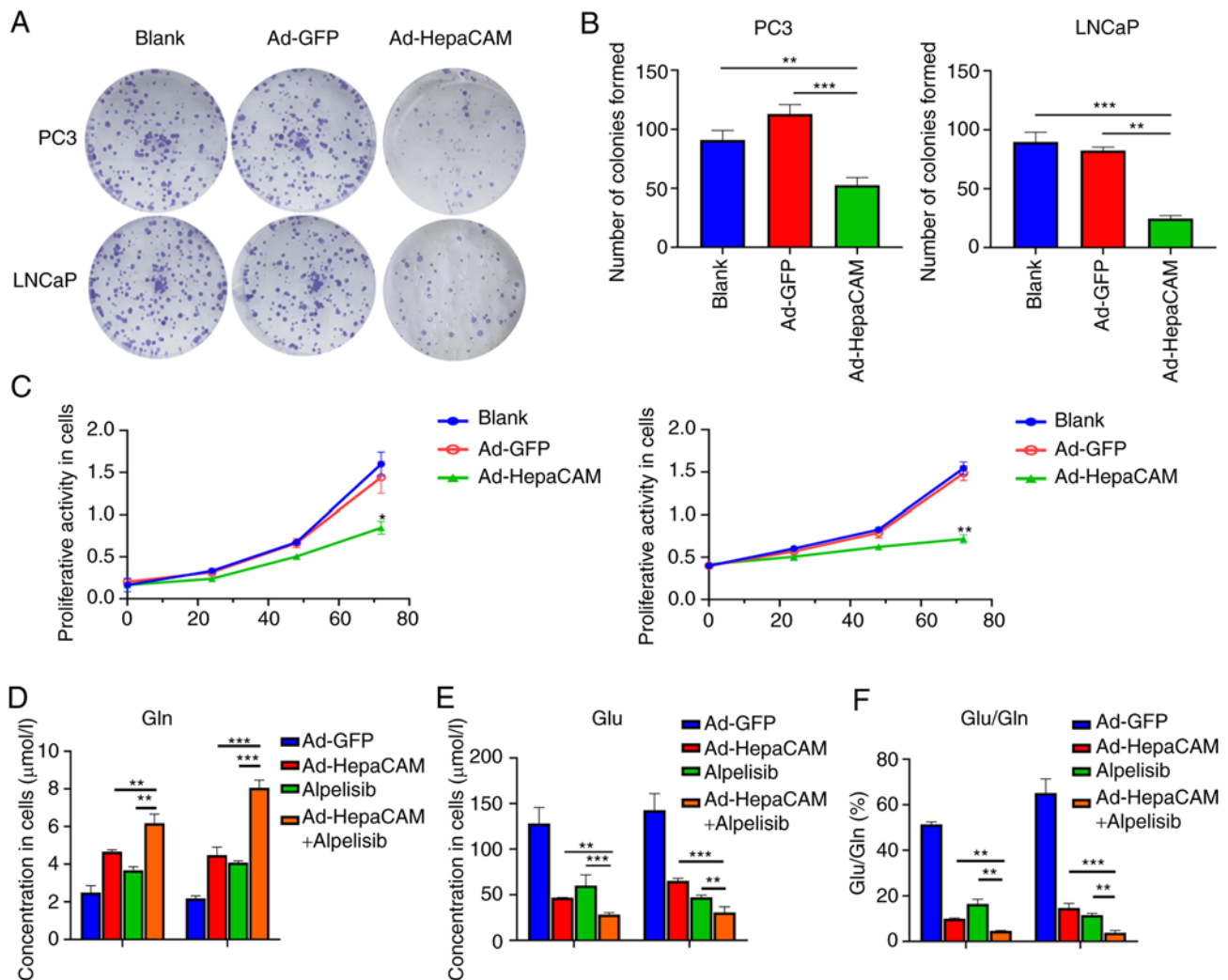


Figure 4. Overexpression of HepaCAM inhibits glutamine metabolism reprogramming and proliferation in PCa cells. (A) Colony formation assay and (B) statistical analysis of the numbers of colonies in the cells after the treatment of Ad-HepaCAM. (C) The proliferative activity in cells after the treatment of Ad-HepaCAM by CCK-8 assay. (D-F) Mass spectrometry tested the concentration of (D) Gln and (E) Glu in the PCa cells after the treatment of Ad-HepaCAM or/and alpelisib, and (F) comparison of the ratio of Glu to Gln. Data are represented as mean \pm SD of three individual experiments. * $P < 0.05$, ** $P < 0.01$, and *** $P < 0.001$ vs. controls. HepaCAM, hepatocellular cell adhesion molecule; PCa, prostate cancer; Gln, glutamine; Glu, glutamate.

that metabolic changes in Gln occurred in patients with PCa, which indicated the potential clinical significance of using Gln as a marker for clinical PCa screening.

High negative correlation between HepaCAM and PIK3CA expression in clinical PCa samples. The expression levels of HepaCAM and PIK3CA in 17 BPH and 44 PCa tissue samples were measured using IHC staining. The results showed that the positive expression rate of HepaCAM reached 34.09% (15/44) among the PCa samples, compared with 76.47% (13/17) in the BPH samples (Fig. 2I). In addition, the positive expression rate of PIK3CA was 86.36% (38/44) in PCa compared with 64.71% (11/17) in the BPH samples (Fig. 2J). Subsequently, HepaCAM and PIK3CA expression were found to be significantly correlated with the Gleason grade (Fig. 2E-H; Table II). Semi-quantitative staining scores revealed significantly decreased HepaCAM expression ($P < 0.05$; Fig. 2I) and significantly increased PIK3CA expression ($P < 0.01$; Fig. 2J) in the PCa tissues compared with those in the BPH tissues. Spearman's correlation analysis calculated that the expression

of HepaCAM was negatively correlated with that of PIK3CA ($P < 0.0001$; Fig. 2K).

Overexpression of HepaCAM inhibits the expression of PIK3CA in PCa cells. The expression of HepaCAM and PIK3CA in normal prostate epithelial cells (RWPE-1) compared with PCa cell lines PC3 and LNCaP were next compared *in vitro*. As shown in Fig. 3A and B, both mRNA and protein expression levels of HepaCAM in the cancer cell lines were significantly lower compared with those in the RWPE-1 cells, while the expression levels of PIK3CA were significantly higher in the cancer cell lines ($P < 0.05$). To ultimately establish the role of HepaCAM in Gln metabolism and proliferation in the PCa cells, Ad-HepaCAM was first transfected into the PC3 and LNCaP cells. The results revealed significantly increased expression levels of HepaCAM in the PC3 and LNCaP cells in the Ad-HepaCAM group compared with those in the Ad-GFP group at both the mRNA and protein levels ($P < 0.001$; Fig. 3C-E). In addition, the expression of PIK3CA in both PC3 and LNCaP cells was significantly

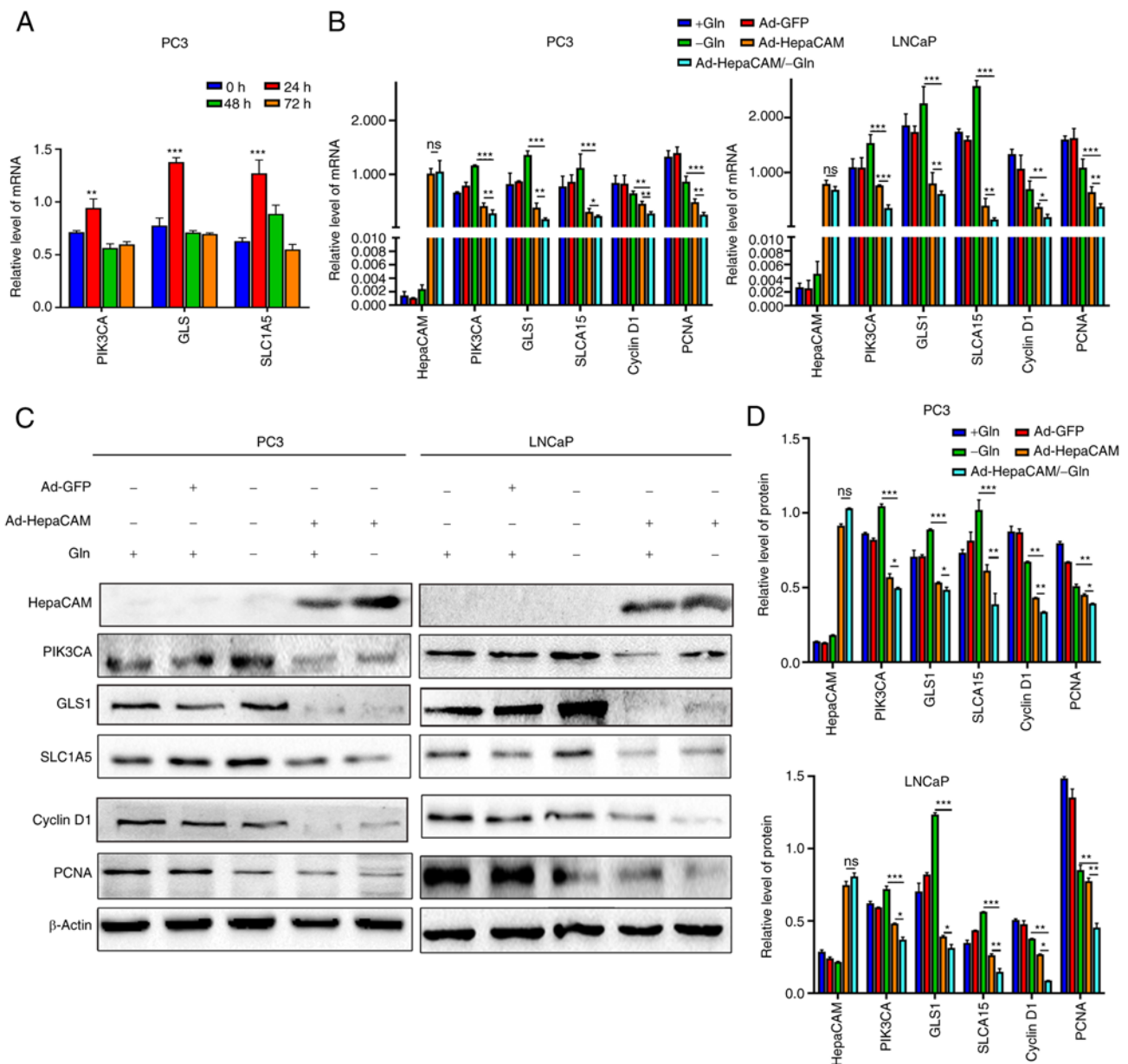


Figure 5. Overexpression of HepaCAM reverses stress resistance to reprogramming of glutamine metabolism in PCa cells. (A) mRNA expression levels of PIK3CA, GLS and SLC1A5 were measured at different time points after Gln deprivation by RT-qPCR. ** $P < 0.01$, and *** $P < 0.001$ vs. 0 h. (B) RT-qPCR detection of mRNA levels of HepaCAM, PIK3CA, GLS, SLC1A5, Cyclin D1, and PCNA in PC3 and LNCaP cells after Ad-HepaCAM or Gln deprivation (24 h). (C and D) Western blot analysis and quantification of the protein expression of the molecules mentioned in (B). β -actin were used as internal controls. Data are represented as mean \pm SD of three individual experiments. * $P < 0.05$, ** $P < 0.01$, and *** $P < 0.001$; ns, not significant. HepaCAM, hepatocellular cell adhesion molecule; PIK3CA, phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit α ; GLS, glutaminase; SLC1A5, solute carrier family 1 member 5; PCNA, proliferating cell nuclear antigen.

downregulated by the overexpression of HepaCAM ($P < 0.01$; Fig. 3C-E).

Overexpression of HepaCAM inhibits Gln metabolism reprogramming and proliferation in PCa cells. Subsequently, the present study next investigated whether overexpression of HepaCAM can mediate an impact on Gln metabolism reprogramming and PCa cell proliferation. The results revealed that compared with that in the blank and Ad-GFP groups, the expression in PC3 and LNCaP cells of Gln metabolism components GLS and SLC1A5 in addition to cyclin D1 and PCNA, proteins associated with proliferation, were significantly reduced in the Ad-HepaCAM group at the mRNA and protein

levels ($P < 0.05$; Fig. 3F-H). Consistently, the results showed that the overexpression of HepaCAM also significantly inhibited the proliferation of PCa cells according to colony formation and CCK-8 assays compared with those in the Ad-GFP group ($P < 0.05$; Fig. 4A-C). MS analysis in PCa cells revealed that Gln (Fig. 4D) levels in the Ad-HepaCAM group were higher compared with those in the Ad-GFP group, while the levels of Glu were lower (Fig. 4E), rendering the ratio of Glu/Gln to be lower as a result of the overexpression of HepaCAM (Fig. 4F). This suggest that Gln metabolism reprogramming was inhibited. These results suggest that HepaCAM overexpression can inhibit Gln metabolism reprogramming and proliferation in PCa cells.

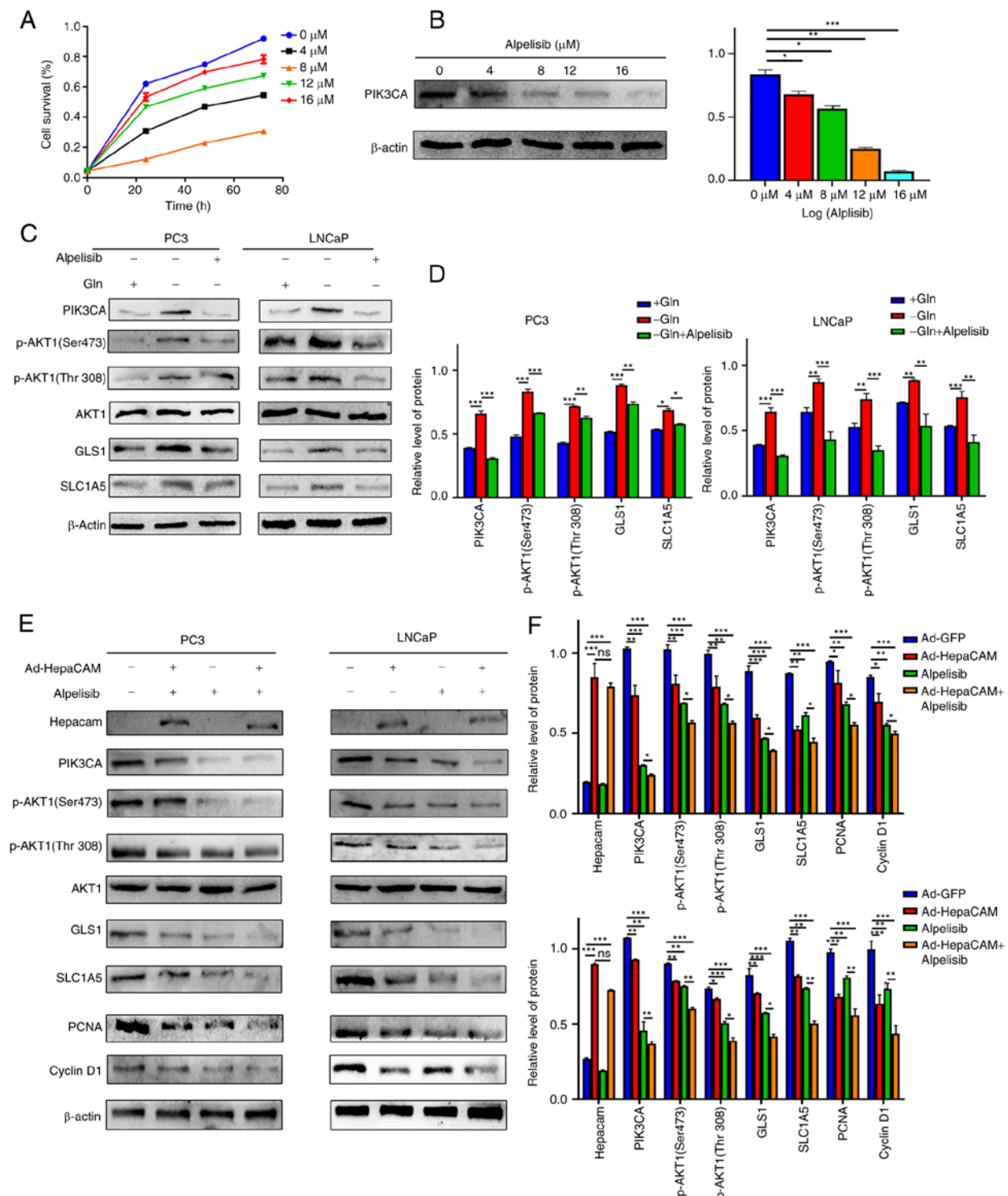


Figure 6. HepaCAM suppresses expression of glutamine metabolism-related and proliferation-related molecules by modulating PIK3CA. (A) CCK-8 assay was used to detect the toxicity of different concentrations of alpelisib (0, 4, 8, 12 and 16 μ M) on PC3 cells. (B) Western blot analysis was used to assess the inhibitory effect on PIK3CA expression with the concentrations of alpelisib used in A. (C and D) Western blot analysis was used to detect and analyse the protein expression of PIK3CA, p-AKT1(Ser473/Thr308), AKT1, GLS1, SLC1A5 after Gln deprivation or/and alpelisib treatment in PC3 and LNCaP cells (8 μ M, 24 h). (E and F) Western blot analysis was used to detect and analyse the protein expression of HepaCAM, cyclin D1, PCNA and the molecules mentioned in C after Gln deprivation for 24, then treatment with Ad-HepaCAM or/and alpelisib for 72 h. β -actin was used as internal controls. Data are represented as mean \pm SD of three individual experiments. * P <0.05, ** P <0.01, and *** P <0.001 vs. controls. HepaCAM, hepatocellular cell adhesion molecule; PIK3CA, phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit α ; GLS, glutaminase; SLC1A5, solute carrier family 1 member 5; PCNA, proliferating cell nuclear antigen; Gln, glutamine.

Stress resistance by Gln metabolism can be reversed by HepaCAM overexpression. Despite the clear inhibitory effects on cell proliferation incurred by limited Gln levels,

cancer cells have the capacity to adapt to the conditions to promote survival and sustain proliferative functions (42). Measuring the expression of enzymes associated with

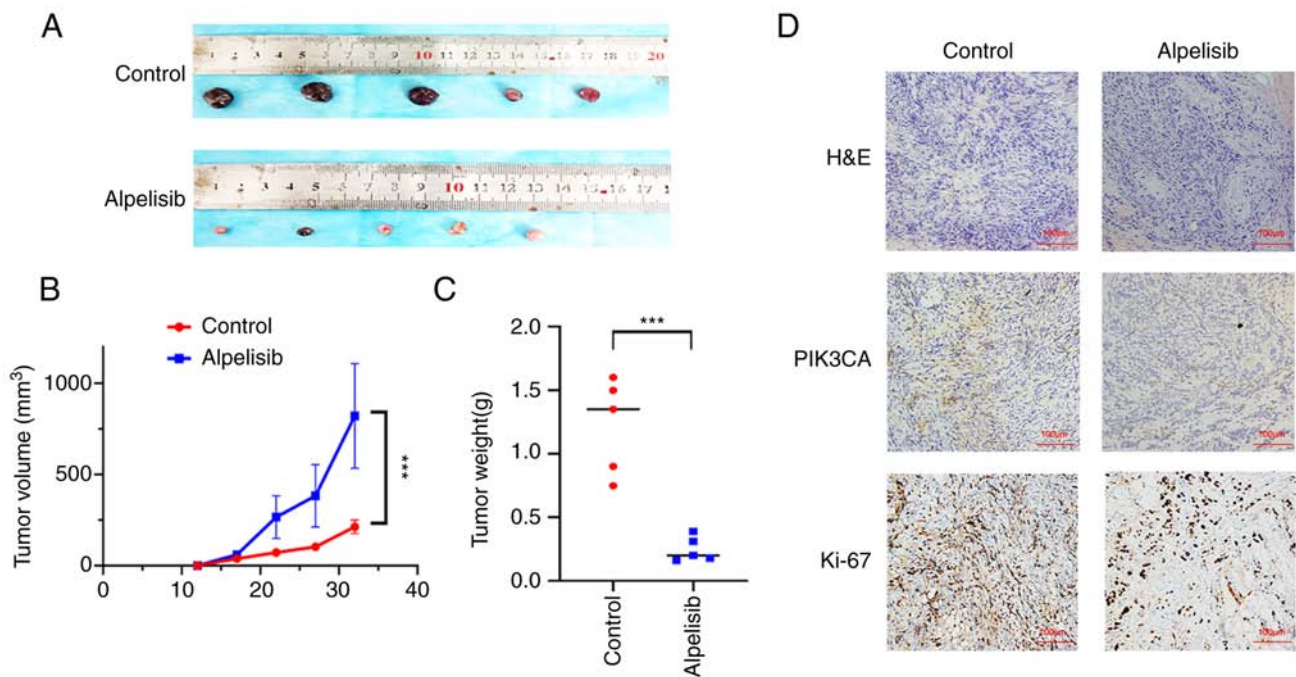


Figure 7. (A-C) Alpelisib significantly suppressed xenograft tumour growth *in vivo*. (D) IHC staining for the expression of PIK3CA and Ki67 in tumour tissue in the alpelisib group and matched group. Magnification, x100. PIK3CA, phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit α . *** $P < 0.001$ vs. controls.

metabolism revealed that the expression of PIK3CA, GLS and SLC1A5 were significantly upregulated immediately after Gln deprivation for 24 h ($P < 0.05$; Fig. 5A). Next, to explore the effects of HepaCAM on this positive response to nutrient starvation in PCa cells, particularly to Gln depletion, Ad-HepaCAM transfection was combined with Gln deprivation for 24 h alongside the corresponding control groups. RT-qPCR and western blotting analyses found that the overexpression of HepaCAM could reverse the upregulation of PIK3CA, GLS and SLC1A5 expression after Gln deprivation. Simultaneously, HepaCAM overexpression was found to significantly reduce the expression of proliferation-associated proteins cyclin D1 and PCNA ($P < 0.05$; Fig. 5B-D). These results suggest that the overexpression of HepaCAM can reverse the metabolic stress resistance of PCa cells after Gln deprivation.

Stress resistance of Gln metabolism is mediated by PIK3CA. To ascertain whether the stress resistance demonstrated by Gln metabolism in PCa cells in response to a poor nutritional environment is associated with PIK3CA, alpelisib was used to suppress and inhibit PIK3CA expression. The minimum inhibitory concentration of cell viability exerted by alpelisib (8 μ M) was first determined using CCK-8 assay ($P < 0.05$; Fig. 6A), while the trend of dose-independent inhibition was demonstrated according to western blotting results ($P < 0.05$; Fig. 6B). It was subsequently discovered that the stress resistance of Gln metabolism in PCa cells was reversed after PIK3CA expression was inhibited by alpelisib under Gln deprivation ($P < 0.05$; Fig. 6C and D). To verify the direct potentiating effects of PIK3CA on maintaining the stress resistance of Gln metabolism *in vivo*, LNCaP cells were subcutaneously injected into nude mice. In total, five mice were subcutaneously injected with alpelisib before the extent of tumour

growth was measured over time. Alpelisib in the dosing group was found to significantly inhibit the growth and reduce the weight of xenograft tumours compared with those in the control group ($P < 0.001$; Fig. 7A-C). In addition, the expression levels of PIK3CA and Ki-67 in the tumour tissues of mice were suppressed by alpelisib (Fig. 7D). In conclusion, these results suggest that PIK3CA can mediate the stress resistance of Gln metabolism in PCa cells.

HepaCAM can suppress Gln metabolism reprogramming and cell proliferation of PCa by inhibiting PIK3CA expression. The present study next examined whether HepaCAM can regulate Gln metabolism reprogramming in PCa by regulating PIK3CA expression. To evaluate this hypothesis, Gln-deprived cells were treated with alpelisib and/or transfected with Ad-HepaCAM. The results demonstrated that the levels of PIK3CA protein expression and its downstream effector p-AKT1 (Ser473/Thr308), Gln metabolism components GLS1 and SLC1A5, proliferation-associated molecules cyclin D1 and PCNA were all significantly lower in the Ad-HepaCAM or alpelisib groups compared with those in their corresponding control groups. In particular, the inhibitory effects mediated by HepaCAM overexpression combined with alpelisib treatment were the most potent ($P < 0.05$; Fig. 6E and F). In addition, suppression of PIK3CA expression did not exert any significant effects on HepaCAM expression (Fig. 6E and F).

Next, CCK-8 assays were performed to detect the sensitivity of PC3 and LNCaP cells to alpelisib after Ad-HepaCAM transfection. The data revealed that alpelisib sensitivity was significantly increased in both cell lines after the overexpression of HepaCAM compared with that in the Ad-GFP group ($P < 0.05$, Fig. 8A). Furthermore, the results showed that the positive expression index of Ki-67, a marker of cancer cell proliferation activity, was significantly

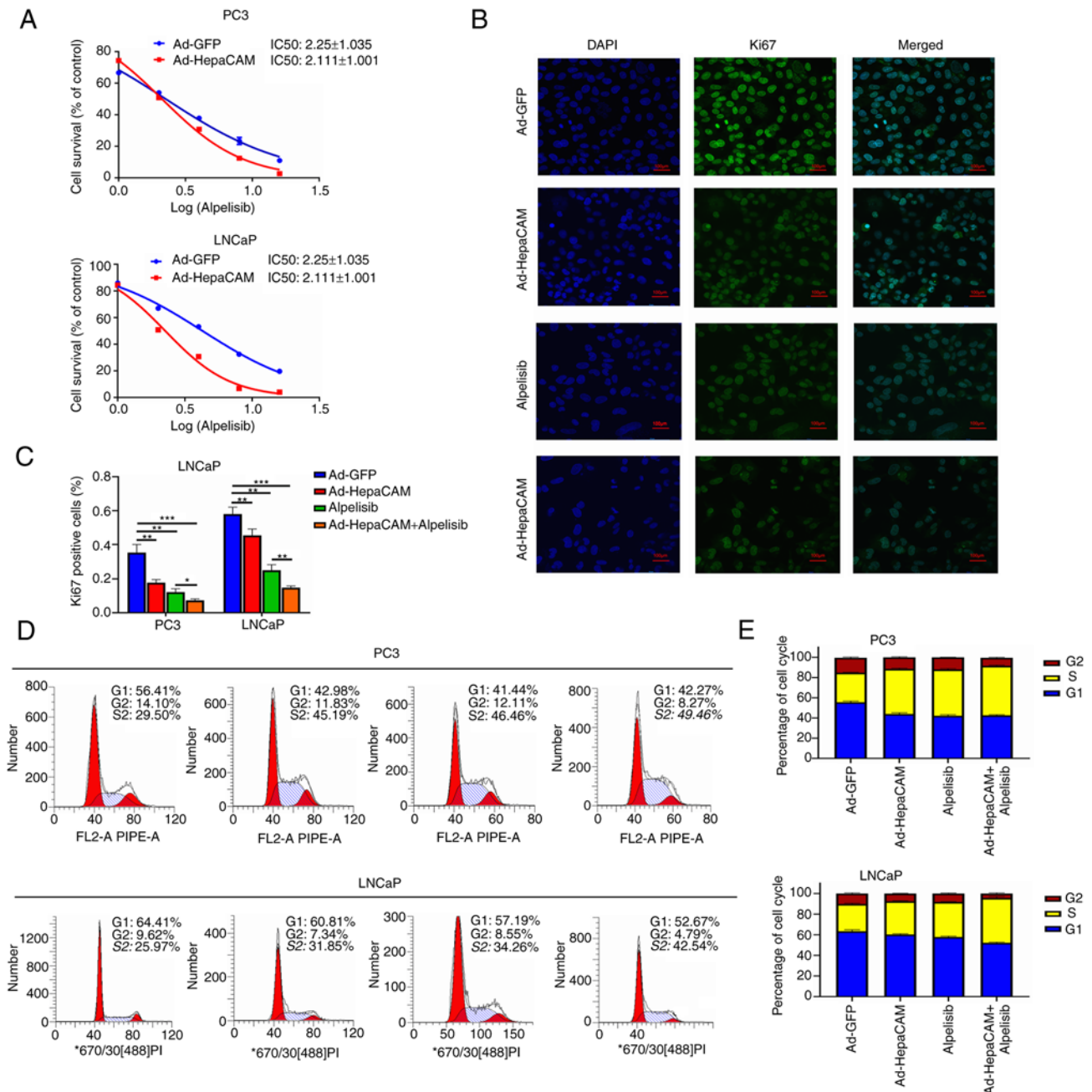


Figure 8. HepaCAM regulates glutamine metabolism reprogramming and cell proliferation of PCa by modulating PIK3CA. (A) IC₅₀ values of alpelisib in PC3 and LNCaP cells were measured by CCK-8 assay in Ad-GFP or Ad-HepaCAM groups (450 nm, 72 h). (B) Ki-67 nuclear expression and (C) positive index in PC3 and LNCaP cells were detected by IF after the treatment of Ad-HepaCAM or/and alpelisib. *P<0.05, **P<0.01, and ***P<0.001. (D) Flow cytometry detected cell cycle distribution after cells were cultured with Ad-HepaCAM or/and alpelisib, and (E) statistical analysis. Data are represented as mean ± SD of three individual experiments. HepaCAM, hepatocellular cell adhesion molecule; PCa, prostate cancer; IC₅₀, half maximal inhibitory concentration.

lower compared with that in the Ad-GFP group following Ad-HepaCAM transfection and/or alpelisib, where the effects exerted by the combination of both treatments were also satisfactory (P<0.05; Fig. 8B and C). Flow cytometry assay revealed that the proportion of cells in the S phase was increased after transfection with Ad-HepaCAM and alpelisib (Fig. 8D and E). By contrast, MS results revealed that the combination of alpelisib and HepaCAM overexpression increased the concentration of Gln further in the PCa cells while suppressing its conversion to Glu (P<0.05; Fig. 4D-F). These results suggest that HepaCAM overexpression can inhibit Gln metabolic reprogramming in PCa cells through

PIK3CA to reverse their stress resistance, which in turn inhibits cell proliferation and tumour growth.

Discussion

In addition to essential sugars and fatty acids, glutamine (Gln) is another important source of nutrients that can accelerate the mitochondrial Krebs cycle, in a process known as glutaminolysis, which involves the catabolism of Gln into glutamate (Glu), which is then metabolized into α-ketoglutarate to support ATP production by feeding into the Krebs cycle (5). In certain cases, cancer cells can reprogram themselves to

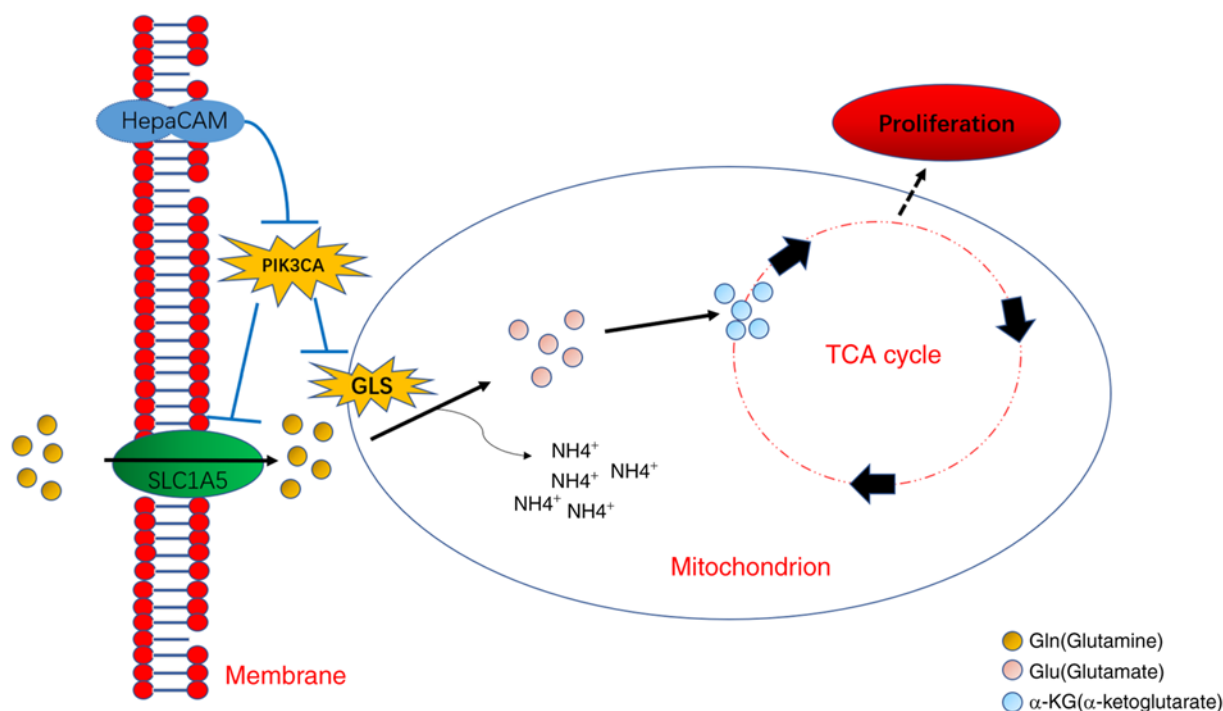


Figure 9. The mechanism of HepaCAM in regards to the inhibition of glutamine metabolism of PCa. HepaCAM inhibits glutamine metabolism-related molecules GLS1, SLC1A5 expression by PIK3CA signalling and reduces transportation and decomposition of glutamine, thereby inhibiting the proliferation driven by the tricarboxylic acid cycle (TCA) cycle. HepaCAM, hepatocellular cell adhesion molecule; PCa, prostate cancer; PIK3CA, phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit α ; GLS, glutaminase; SLC1A5, solute carrier family 1 member 5.

become dependent on certain metabolic pathways, activities or processes, such that the cell adopts a type of oncogenic addiction, in a phenomenon known as ‘metabolic reprogramming’ (6). Importantly, this type of metabolic reprogramming can be tissue-specific, and in context, downstream of both genetic and environmental factors, such that it can present vulnerabilities that can be therapeutically exploited (43,44). Therefore, the present study compared the levels of various amino acids in the blood samples of normal individuals and patients with prostate cancer (PCa). Gln levels exhibited one of the most sensitive and significant differences, which was in turn correlated with the Gleason score based on the clinical parameters of the patients. In addition, through the detection and analysis of the levels of Gln and metabolic flux in the blood samples and PCa cells, it was found that Gln metabolism reprogramming was also common among patients with PCa. Therefore, it was hypothesized that Gln levels have potential as a type of auxiliary diagnostic marker for PCa in a clinical setting.

Previous studies reported that hepatocellular cell adhesion molecule (HepaCAM) can serve a significant role in tumour cell proliferation, apoptosis, migration and invasion, since its expression is typically absent in cancer tissues (19,45). In addition, previous reports have found that mutation in phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit α (PIK3CA) can activate PI3K/AKT signalling downstream to induce alterations in Gln metabolism in tumour cells (29). In the present study, bioinformatics analyses of various databases were performed to predict the correlation in the expression levels between HepaCAM and PIK3CA. These data revealed that PIK3CA and that of glutaminase (GLS) and as solute

carrier family 1 member 5 (SLC1A5), components of Gln metabolism, are expressed at high levels in PCa tissues, all of which in turn correspond to poorer prognoses. In addition, a negative correlation was found between HepaCAM and PIK3CA expression in PCa tissues as determined by IHC. Subsequently, HepaCAM and PIK3CA expression were found to be significantly correlated with the Gleason grade, which implies the different prognoses associated with their expression levels in PCa tissues. Therefore, these findings suggest that HepaCAM can regulate Gln metabolic reprogramming in PCa by regulating PIK3CA expression.

At present, the most extensively studied metabolic alteration in cancer cells is their high reliance on glycolysis even under sufficient oxygen supply to sustain oxidative phosphorylation in the mitochondria, in a process known as the Warburg effect (19). The rapid ATP production as a result of this glycolytic upregulation is considered to potentiate the proliferation of cancer cells (46). However, it has been previously reported that the catabolic and anabolic roles of both glucose and Gln can cooperate in promoting tumour growth (47). A previous study found that HepaCAM can inhibit the Warburg effect in renal cell carcinoma cells (19). Therefore, it could be hypothesized that HepaCAM can inhibit Gln metabolism reprogramming in PCa. In the present study, the mRNA and protein expression levels of HepaCAM and PIK3CA in normal prostate epithelial cells RWPE-1 and prostate cancer cell lines PC3 and LNCaP were next assessed. After the induced overexpression of HepaCAM in the cells lines, the expression of PIK3CA and Gln metabolism components GLS and SLC1A5, in addition to tumour proliferation-associated molecules PCNA and cyclin D1, were all reduced. These results suggest that the loss of HepaCAM expression in tumours may

promote dysfunctions in Gln metabolism, promote cell proliferation and tumour development.

Subsequently, the results from the present study revealed stress resistance or a compensatory effect in Gln metabolism in PCa cells following Gln starvation. GLS and SLC1A5 expression were temporarily increased in the short-term after Gln deprivation in cell culture conditions, which may be a potential cause of the ineffectiveness of Gln starvation in PCa. On this basis, HepaCAM was overexpressed, which eliminated this stress compensation effect. In addition, the inhibitory effects of HepaCAM combined with Gln deprivation on tumour proliferation were significantly more potent compared with those mediated by either treatment alone. Although this compensatory response has been previously reported in cervical cancer tissues, a previous study found this effect after treatment with inhibitors of GLS1 (48). In addition, another previous study discovered that arginine can participate in the mediation of this type of compensation (49). Since abnormal arginine levels were also found in the blood samples of patients with PCa in the present study, the possibility of other signalling pathways mediating this type of stress compensation apart from HepaCAM cannot be ruled out.

The present study then explored the effects of HepaCAM on Gln metabolism reprogramming of PCa. As one of the catalytic subunits of classical PI3K, PIK3CA serves a key role in activating the PI3K/AKT signalling pathway (25). A number of studies have previously found that PIK3CA has a high frequency of mutations in tumours (40,41,50). Through bioinformatics analysis, it was found that PIK3CA primarily serves a facilitating role in PCa pathophysiology, where it was frequently found to be activated by amplification. This suggests that this form of PIK3CA activation is tissue-specific. Subsequently, higher PIK3CA expression levels were also found in PCa tissues compared with those in the adjacent normal tissues, where they in turn were correlated inversely and significantly with those of HepaCAM. Alpelisib is a specific inhibitor of PIK3CA (51). The present study found that inhibition of PIK3CA expression by alpelisib significantly reversed the stress resistance of PCa cells to Gln deprivation, supporting the possible effects of PIK3CA in this phenomenon. After the PCa cells deprived of Gln for 24 h were treated with alpelisib and transfected with Ad-HepaCAM at the same time, the phosphorylation of AKT, the expression of Gln metabolism-associated and proliferation-associated proteins were all reduced. In addition, the suppression of PIK3CA exerted no significant impact on HepaCAM expression. These observations suggest that HepaCAM can strengthen the inhibition of Gln metabolic reprogramming and proliferation in PCa by reversing stress resistance through inhibiting PIK3CA expression. However, the present study did not assess the possibility of a direct interaction between HepaCAM and PIK3CA at the protein level. Instead, the protein structure database STRING was used to predict protein interaction, showing that HepaCAM may bind to PIK3CA directly. However, this requires further experimental verification.

Alpelisib is a PIK3CA inhibitor that has demonstrated antitumor activity in several types of cancer, including breast cancer (52) and epithelial ovarian cancer (53). In addition, it has been previously applied in clinical trials for the treatment of multiple tumours. Results from the present study suggest that it exerts significant inhibitory effects on PIK3CA in PCa

cells in a dose-dependent manner. Although HepaCAM was shown to inhibit Gln metabolism reprogramming and proliferation in PCa, the addition of alpelisib was able to significantly potentiate the inhibition caused by HepaCAM overexpression. Although a specific agonist of HepaCAM has not yet been successfully developed, the use of PIK3CA inhibitors to block tumour growth and cell proliferation appear to more likely result in greater clinical significance, which provides a valuable reference for the clinical treatment of PCa.

The present study has a number of limitations. Although the mRNA expression data of HepaCAM in the bioinformatics analysis is the relative expression value obtained after standardization according to the FPKM algorithm, the abundance value of HepaCAM expression in the TCGA PCa datasets were relatively low, which could have caused a certain degree of interference in the follow-up IHC experimental conclusions.

In summary, the present study uncovered abnormal Gln concentrations, which may be exploited as a potential indicator for the diagnosis of PCa. In addition, to the best of our knowledge, the present study was the first to discover that the HepaCAM/PIK3CA axis may be implicated in Gln metabolism reprogramming and PCa proliferation (Fig. 9). Mechanistically, HepaCAM overexpression reversed the stress resistance shown by Gln metabolism by downregulating PIK3CA expression. Importantly, the combination of PIK3CA inhibitor and overexpression of HepaCAM was able to effectively inhibit Gln metabolism reprogramming and PCa proliferation, thereby providing a certain clinical value for the treatment of PCa.

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Availability of data and materials

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

ZH, CL and YG designed the experiments. ZH, YG, TL and CY collected the specimens and analysed the clinical data. ZH, YG, TL and CY performed the experiments. CL and LO provided technical support in this research project and supervised the progress of the experiments. ZH and YG analysed the statistical data. ZH, LO and TL assembled and installed the figures. ZH, YG and TL confirm the authenticity of all the raw data. All authors have read and approved the final manuscript for publication.

Ethics approval and consent to participate

The present study was approved by the Ethics Committee of Chongqing Medical University (2016.12.16 Chongqing, China). Informed consent was obtained from the patients or their family members who agreed to the use of their samples in this study. The animal study was performed in accordance with the Animal Research Advisory Committee (ARAC) Guidelines and was approved by the Ethics Committee of Chongqing Medical University (Chongqing, China).

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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